

**Stantec Analytical Validation Checklist****Report No. ATA37**

Project Name: Amtrak North Yard	Project Number: 213402048	
Validator: Jim Tezak	Laboratory: Eurofins/Lancaster Laboratory	
Date Validated: 12/12/2019	Laboratory Project Number: 1595161	
Sample Start-End Date: 9/22/2015	Laboratory Report Date: 9/30/2015	
Parameters Validated: Polychlorinated biphenyls (PCBs) by EPA SW-846 3580A/8082A - oil matrix		
Samples Validated (all Grab Soil): MH-14 Area-Oil, LLI # 8059389		
<b>VALIDATION CRITERIA CHECK</b>		
Validation Flags Applicable to this Review: <b>U</b> The analyte was analyzed for, but not detected above the reported sample quantitation limit. <b>J</b> The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample. <b>J+</b> Result is estimated quantity but the result may be biased high. <b>J-</b> Result is estimated quantity but the result may be biased low. <b>UJ</b> The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample. <b>NJ</b> The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated numerical value represents its approximate concentration. <b>B</b> The analyte was detected in the method, field, and/or trip blank. <b>R</b> The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.		
1. Were all the analyses requested for the samples submitted with each COC completed by the lab?	Yes <b>X</b>	No
Comments:		
2. Did the laboratory identify any non-conformances related to the analytical result?	Yes	No <b>X</b>
Comments:		
3. Were sample Chain-of-Custody forms complete?	Yes <b>X</b>	No
Comments: Samples were listed on chain-of-custody (COC) # 194670.		
4. Were samples received in good condition and at the appropriate temperature?	Yes <b>X</b>	No
Comments: The laboratory noted on the Sample Administration Receipt Documentation Log that the shipping container was not sealed and there was no custody seal present when the samples were received.		

5. Were sample holding times met?		Yes <b>X</b>	No
Comments:			
6. Were correct concentration units reported?		Yes <b>X</b>	No
Comments: Results were reported in units of microgram per kilogram (ug/kg).			
7. Were detections found in laboratory blank samples?		Yes	No <b>X</b>
Comments:			
8. Were detections found in field blank, equipment rinse blank, and/or trip blank samples?	NA <b>X</b>	Yes	No
Comments: No field blanks were submitted in this sample delivery group (SDG).			
9. Were instrument calibrations within method criteria?	NA <b>X</b>	Yes	No
Comments: Not Applicable, Level 2 data validation.			
10. Were surrogate recoveries within control limits?		Yes <b>X</b>	No
Comments:			
11. Were laboratory control sample(s) (LCS/LCSD) sample recoveries within control limits?		Yes <b>X</b>	No
Comments:			
12. Were matrix spike (MS/MSD) recoveries within control limits?	NA <b>X</b>	Yes	No
Comments: Not applicable; site-specific MS/MSD not analyzed for this SDG.			
13. Were RPDs within control limits?		Yes <b>X</b>	No
Comments: Site-specific MS/MSD not analyzed. The laboratory reported LCS/LCSD results to assess accuracy and precision. All LCS/LCSD recoveries and RPDs were within control limits.			
14. Were dilutions required on any samples?		Yes <b>X</b>	No
Comments: The sample was diluted 20X prior to analysis. Sample reporting limits were adjusted accordingly. No data were qualified.			

15. Were Tentatively Identified Compounds (TIC) present?	NA <b>X</b>	Yes	No
Comments: TIC not requested.			
16. Were organic system performance criteria met?	NA <b>X</b>	Yes	No
Comments: Not Applicable, Level II data validation.			
17. Were GC/MS internal standards within method criteria?	NA <b>X</b>	Yes	No
Comments: Not Applicable, Level II data validation.			
18. Were inorganic system performance criteria met?	NA <b>X</b>	Yes	No
Comments:			
19. Were blind field duplicates collected? If so, discuss the precision (RPD) of the results.		Yes	No <b>X</b>
<b><u>Duplicate Sample ID</u></b>		<b><u>Primary Sample No.</u></b>	
Comments: No PCB Aroclors were detected in either sample.			
20. Were at least 10 percent of the hard copy results compared to the Electronic Data Deliverable Results?	Yes	No <b>X</b>	Initials JET
Comments: At the time data verification was performed, electronic data had not been loaded into the project database, so the comparison of hard copy results to EDD results could not be completed. After the data are loaded into the database, a review of hard copy results versus the electronic data will be performed.			
21. Other?		Yes	No <b>X</b>
Comments: All samples were validated according to the USEPA 2014 NFGs and DNREC SOPCAP. All data are considered usable as qualified. No data have been rejected.			
<b>PRECISION, ACCURACY, METHOD COMPLIANCE AND COMPLETENESS ASSESSMENT</b>			
Precision:	Acceptable <b>X</b>	Unacceptable	Initials JET
Comments:			
Sensitivity:	Acceptable <b>X</b>	Unacceptable	Initials JET
Comments:			

Accuracy:	Acceptable X	Unacceptable	Initials JET
Comments:			
Representativeness:	Acceptable X	Unacceptable	Initials JET
Comments:			
Method Compliance:	Acceptable X	Unacceptable	Initials JET
Comments:			
Completeness:	Acceptable X	Unacceptable	Initials JET
Comments:			